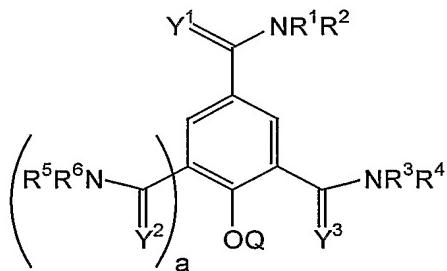


**WHAT IS CLAIMED IS:**

1           **1.**       A compound having the structure:

2



3           wherein

4           R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup> and R<sup>6</sup> are members independently selected from H,  
5           substituted or unsubstituted alkyl, substituted or unsubstituted  
6           heteroalkyl, substituted or unsubstituted aryl, and substituted or  
7           unsubstituted heterocycloalkyl, wherein a member selected from R<sup>1</sup>  
8           and R<sup>2</sup>; R<sup>3</sup> and R<sup>4</sup>; and R<sup>5</sup> and R<sup>6</sup>, together with the nitrogen atom  
9           to which they are attached, optionally form a ring system selected  
10          from heteroaryl and heterocycloalkyl;

11          Y<sup>1</sup>, Y<sup>2</sup> and Y<sup>3</sup> are members independently selected from O and (H)<sub>2</sub>;

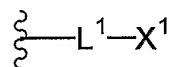
12          Q is a member selected from H, a protecting group and a cleaveable group;

13          and

14          a is 0 or 1.

1           **2.**       The compound according to claim **1**, wherein a member selected  
2          from R<sup>1</sup>, R<sup>3</sup> and R<sup>5</sup> has the structure:

3

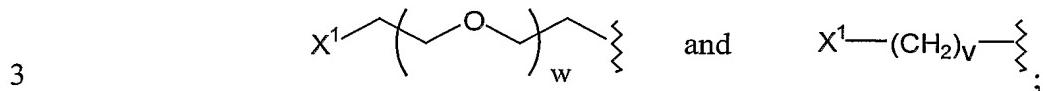


4           wherein

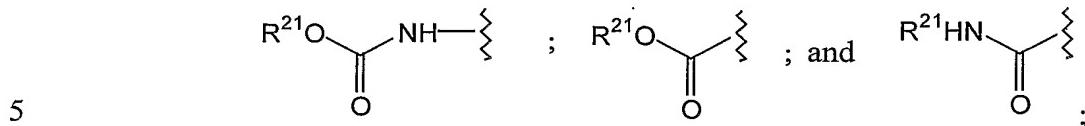
5           L<sup>1</sup> is a member selected from substituted or unsubstituted alkyl, substituted  
6           or unsubstituted heteroalkyl and substituted or unsubstituted aryl;  
7           and

8           X<sup>1</sup> is a member selected from protected or unprotected reactive functional  
9           groups and non-covalent protein binding groups.

1           **3.**       The compound according to claim **2**, wherein a member selected  
2          from R<sup>1</sup>, R<sup>3</sup> and R<sup>5</sup> is a member selected from:



4                     $X^1$  is a member selected from:



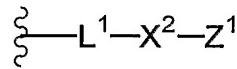
6                    in which  $\text{R}^{21}$  is a member selected from H, substituted or unsubstituted  
7                    alkyl and substituted or unsubstituted aryl;

8                     $v$  is an integer from 1 to 20; and

9                     $w$  is an integer from 1 to 1,000.

1                  4. The compound according to claim 2, wherein said non-covalent  
2                   protein binding group is sulfonate.

1                  5. The compound according to claim 1, wherein a member selected  
2                   from  $\text{R}^1$ ,  $\text{R}^3$  and  $\text{R}^5$  has the structure:



4                    wherein

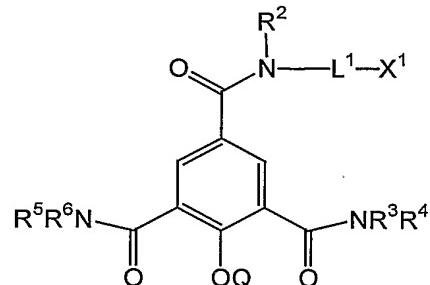
5                     $\text{L}^1$  is a member selected from substituted or unsubstituted alkyl and  
6                    substituted or unsubstituted heteroalkyl; and

7                     $\text{X}^2$  is a linking member adjoining  $\text{L}^1$  to  $\text{Z}^1$ ; and

8                     $\text{Z}^1$  is a member selected from carrier molecules and detectable labels.

1                  6. The compound according to claim 5, wherein said carrier molecule  
2                   is a targeting agent.

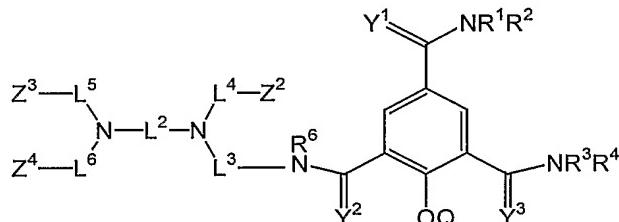
1                  7. The compound according to claim 2, having the structure:



2                    wherein

4            $X^1$  is a member selected from  $\text{NH}_2$ ,  $\text{SH}$ ,  $\text{COR}^7$ ,  $\text{O}(\text{CH}_2)_m\text{Z}^6$ ,  $\text{NHNH}_2$  and  
 5            $\text{O}(\text{CH}_2)_2(\text{OCH}_2\text{CH}_2)_s\text{O}(\text{CH}_2)_2\text{Z}^6$   
 6           wherein  
 7            $R^7$  is a member selected from  $\text{H}$ ,  $\text{OR}^8$ ,  $\text{OCOR}^8$ ,  $\text{NR}^8\text{R}^9$ ,  
 8           wherein  
 9            $R^8$  and  $R^9$  are members independently selected from  $\text{H}$ ,  
 10           substituted or unsubstituted alkyl, substituted or  
 11           unsubstituted heteroalkyl, substituted or  
 12           unsubstituted aryl, substituted or unsubstituted  
 13           heteroaryl and substituted or unsubstituted  
 14           heterocycloalkyl;  
 15            $Z^6$  is a member selected from  $\text{OR}^{10}$ ,  $\text{OCOR}^{10}$ ,  $\text{NR}^{10}\text{R}^{11}$   
 16           wherein  
 17            $R^{10}$  and  $R^{11}$  are members independently selected from  $\text{H}$ ,  
 18           substituted or unsubstituted alkyl, substituted or  
 19           unsubstituted heteroalkyl, substituted or  
 20           unsubstituted aryl, substituted or unsubstituted  
 21           heteroaryl and substituted or unsubstituted  
 22           heterocycloalkyl;  
 23            $m$  is an integer from 1 to 20; and  
 24            $s$  is an integer from 1 to 1000.

1           **8.**       The compound according to claim 1, having the structure:

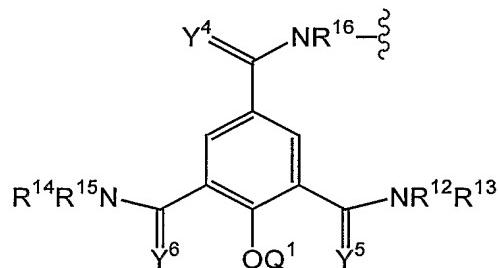


2           wherein

3            $L^2$  is a member selected from substituted or unsubstituted alkyl, substituted  
 4           or unsubstituted heteroalkyl, substituted or unsubstituted aryl,  
 5           substituted or unsubstituted heteroaryl, substituted or unsubstituted  
 6           heterocycloalkyl;  
 7            $L^3$ ,  $L^4$ ,  $L^5$  and  $L^6$  are members independently selected from a single bond,  
 8           substituted or unsubstituted alkyl and substituted or unsubstituted  
 9           heteroalkyl; and  
 10

11            $Z^2$ ,  $Z^3$ , and  $Z^4$  are members independently selected from H, substituted or  
12           unsubstituted aryl and substituted or unsubstituted heteroaryl.

1           **9.**       The compound according to claim 8, wherein  $Z^2$ ,  $Z^3$ , and  $Z^4$  are  
2       members independently selected from substituted or unsubstituted pyridyl, substituted or  
3       unsubstituted salicylamidyl, substituted or unsubstituted phthalamidyl, substituted or  
4       unsubstituted terephthalamidyl, substituted or unsubstituted catechol and

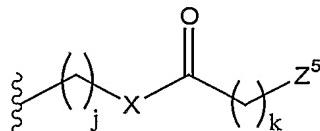


5           wherein

7            $R^{12}$ ,  $R^{13}$ ,  $R^{14}$ ,  $R^{15}$  and  $R^{16}$  are members independently selected from H,  
8       substituted or unsubstituted alkyl, substituted or unsubstituted  
9       heteroalkyl, substituted or unsubstituted aryl, and substituted or  
10      unsubstituted heterocycloalkyl, wherein a member selected from  $R^7$   
11      and  $R^8$ ; and  $R^9$  and  $R^{10}$ , together with the nitrogen atom to which  
12      they are attached, form a ring system selected from heteroaryl and  
13      heterocycloalkyl;  
14       $Y^4$ ,  $Y^5$  and  $Y^6$  are members independently selected from O and (H)2; and  
15      Q is a member selected from H, a protecting group or a cleaveable group.

1           **10.**      The compound according to claim 8, wherein  $L^2$  is a substituted or  
2       unsubstituted C<sub>1</sub>-C<sub>6</sub> alkyl group.

1           **11.**      The compound according to claim 1, wherein at least one of  $R^1$ ,  $R^3$   
2       and  $R^5$  has the structure:

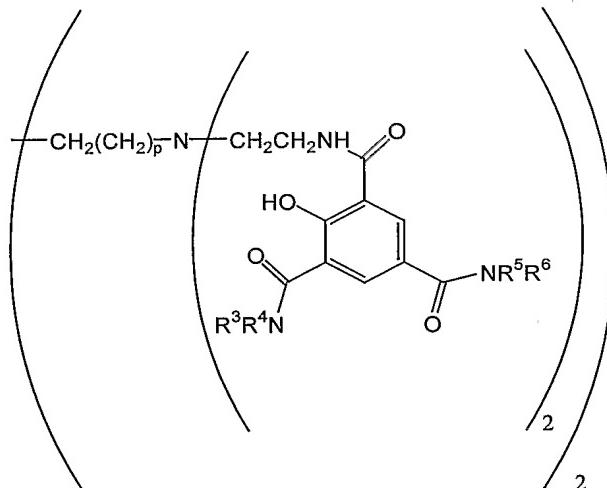


3           wherein,

5            $Z^5$  is a member selected from H,  $OR^{17}$ ,  $SR^{17}$ ,  $NHR^{17}$ ,  $OCOR^{18}$ ,  
6        $OC(O)NHR^{18}$ ,  $NHC(O)OR^{17}$ ,  $OS(O)_2OR^{17}$ , and  $C(O)R^{18}$ ;

7           R<sup>17</sup> is a member selected from H, substituted or unsubstituted alkyl, and  
 8           substituted or unsubstituted heteroalkyl;  
 9           R<sup>18</sup> is a member selected from H, OR<sup>19</sup>, NR<sup>19</sup>NH<sub>2</sub>, SH, C(O)R<sup>19</sup>, NR<sup>19</sup>H,  
 10          substituted or unsubstituted alkyl and substituted or unsubstituted  
 11          heteroalkyl;  
 12          R<sup>19</sup> is a member selected from H, substituted or unsubstituted alkyl and  
 13          substituted or unsubstituted alkyl;  
 14          X is a member selected from O, S and NR<sup>20</sup>  
 15          wherein  
 16           R<sup>20</sup> is a member selected from H, substituted or unsubstituted alkyl  
 17           and substituted or unsubstituted heteroalkyl; and  
 18          j and k are members independently selected from the group consisting of  
 19          integers from 1 to 20.

1           **12.**   The compound according to claim 1, having the structure:

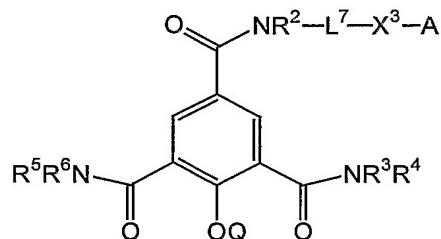


2           in which p is an integer from 0 to 2.

1           **13.**   A polymer comprising a subunit having said structure according to  
 2          claim 1.

1           **14.**   The polymer according to claim 13, wherein said polymer is a  
 2          biomolecule.

1           **15.**   The polymer according to 1, having the structure:



2

3       wherein

4           L<sup>7</sup> is a member selected from a single bond, substituted or unsubstituted  
5           alkyl and substituted or unsubstituted aryl; and6           X<sup>3</sup> is linking member joining L<sup>7</sup> to A;

7           A is a carrier molecule.

1           **16.**   The polymer according to claim **15** wherein A is a member  
2           selected from biopolymers, poly(amino acids), polyethers, polyimines, polysaccharides,  
3           dendrimers, cyclodextrins, pharmaceutical agents.

1           **17.**   The polymer according to claim **16**, wherein said biopolymer is a  
2           member selected from polypeptides, nucleic acids and saccharides.

1           **18.**   The polymer according to claim **17**, wherein said protein is a  
2           member selected from antibodies, enzymes, and serum proteins

1           **19.**   A chelate of a metal ion comprising an organic ligand having said  
2           structure according to claim **1**.

1           **20.**   The chelate according to claim **19**, wherein said metal ion is a  
2           lanthanide ion.

1           **21.**   The chelate according to claim **20**, wherein said chelate is  
2           luminescent.

1           **22.**   The chelate according to claim **19**, wherein said chelate is  
2           covalently attached to a carrier molecule.

1           **23.**   A method for detecting enzyme in a sample, said method  
2           comprising:  
3           (a) contacting said sample with a peptide construct comprising:

**24.** A method of determining the effect of a compound on enzyme activity, said method comprising:

- 3                     (a) contacting a sample comprising said enzyme with a peptide construct  
4                         comprising:  
5                             iii) a peptide sequence, said sequence comprising a cleavage site  
6                                 for said enzyme;  
7                             iv) a complex according to claim 19 covalently bound to said  
8                                 peptide sequence; and  
9                             iii) a quencher of light energy covalently bound to said peptide  
10                                 sequence, said quencher having an absorbance band  
11                                 overlapping an emission band of said complex,  
12                                 wherein said peptide sequence conformation allows light energy  
13                                 transfer between said complex and said quencher when said  
14                                 complex is excited;  
15                     (b) exciting said complex;  
16                     (c) determining a fluorescence property of said sample; and

17                   (d) comparing said fluorescence property from step (c) with a reference  
18                   fluorescence property for said peptide construct, wherein said activity of said  
19                   enzyme in said sample alters said light energy transfer, resulting in a change in  
20                   said fluorescence property.

1                   **25.**       A method for detecting a target nucleic acid sequence, said method  
2                   comprising:

3                   (a) contacting said target sequence with a detector oligonucleotide comprising a  
4                   single-stranded target binding sequence, said detector oligonucleotide having  
5                   covalently linked thereto,

6                   i) a complex according to claim 19;

7                   ii) a quencher of light energy having an absorbance band overlapping  
8                   an emission band of said complex,

9                   wherein said detector nucleic acid conformation allows fluorescence  
10                  energy transfer between said complex and said quencher when said  
11                  complex is excited;

12                  (b) hybridizing said target binding sequence to said target sequence, thereby  
13                  altering said conformation of said detector oligonucleotide, causing a change  
14                  in a fluorescence parameter of said complex; and

15                  (c) determining a fluorescence property of said sample; and

16                  (d) comparing said fluorescence property from step (c) with a reference

17                  fluorescence property for said peptide construct, wherein said activity of said  
18                  enzyme in said sample alters said light energy transfer, resulting in a change in  
19                  said fluorescence property.

1                   **26.**       The method according to claim 25, wherein said detector  
2                  oligonucleotide has a format selected from molecular beacons, scorpion probes, sunrise  
3                  probes, light up probes and TaqMan™ probes.

1                   **27.**       The method according to claim 23, 24 or 25, wherein said  
2                  fluorescence property is detected in-real time.

1                   **28.**       The method according to claim 23, 24 or 25, wherein said change  
2                  and said fluorescence property measured is a change in fluorescence intensity.

1                   **29.**     A microarray comprising a complex according to claim **19**,  
2     wherein said complex is conjugated to a solid support or to a carrier molecule attached to  
3     said solid support.

1                   **30.**     The microarray according to claim **29**, wherein said carrier  
2     molecule is a member selected from a nucleic acid, a peptide, a peptide nucleic acid, a  
3     pharmaceutical agent and combinations thereof.

1                   **31.**     The microarray according to claim **29**, wherein said solid support is  
2     divided into a first region and a second region, said first region having attached thereto a  
3     first complex, and said second region having attached thereto a second.

1                   **32.**     A method of providing radiation therapy to a subject requiring such  
2     therapy, said method comprising:

3                         administering to said subject a complex according to claim **19**, said  
4                         complex having radiosensitization properties; and  
5                         administering ionizing radiation to said subject, thereby providing  
6                         radiation therapy to said subject.

1                   **33.**     A method for photodynamic therapy of a lesion or of a lesion  
2     beneath melanodermic tissue of a subject, said method comprising:

3                         (a) administering a complex according to claim **19** to said subject; and  
4                         (b) photoirradiating said lesion.

1                   **34.**     The method according to claim **33**, wherein said photoirradiating is  
2     with light having a wavelength range of about 610 to about 1150 nanometers.

1                   **35.**     The method of claim **34** wherein the photoirradiating is with light  
2     having a wavelength range of about 730 to about 770 nanometers.

1                   **36.**     The complex according to claim **19**, wherein said complex  
2     comprises a component of an ink or a dye.

1                   **37.**     The complex according to claim **19**, wherein said complex  
2     comprises a component of a substrate for the transmission and amplification of light.

1                   **38.**     The complex according to claim 37, wherein said substrate  
2     comprises a member selected from glass, organic polymers, inorganic polymers and  
3     combinations thereof.

1                   **39.**     A method for amplifying light transmitted by a substrate, said  
2     method comprising transmitting light through a substrate according to claim 37, thereby  
3     amplifying said light.